

In the Claims:

Cancel Claims 1 and 2.

Claims 3 to 7 (cancelled).

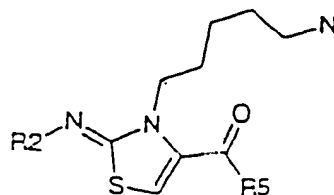
Claim 8. (presently amended) A composition for an affinity for somatostatin receptors comprising an amount of a compound of Claim + 12 sufficient for said affinity and a pharmaceutical carrier.

Claim 9. (presently amended) A method of treating disorders in warm-blooded animals caused by somatostatin receptors comprising administering to warm-blooded animals in need thereof an amount of a compound of Claim + 12 sufficient to treat said disorders.

Cancel Claims 10 and 11.

Add the following Claims :

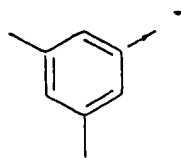
Claim 12. A compound of the formula:



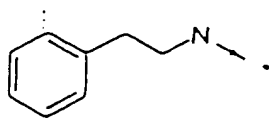
(i)

wherein

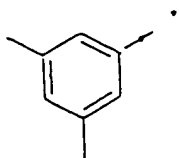
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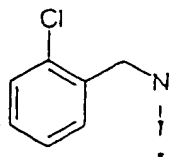
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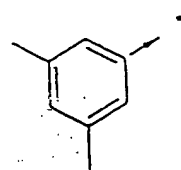
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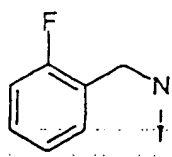
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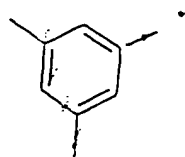
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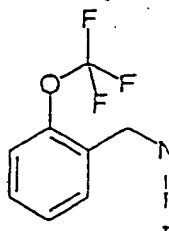
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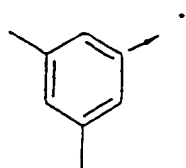
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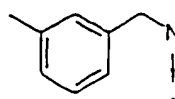
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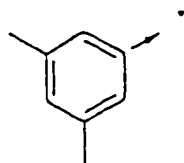
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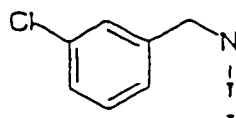
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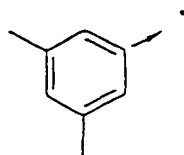
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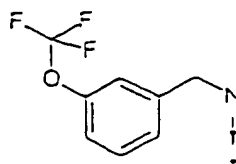
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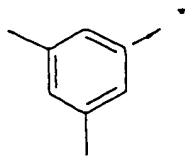
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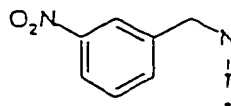
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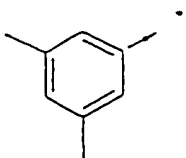
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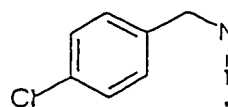
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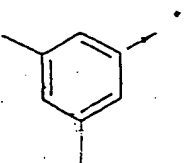
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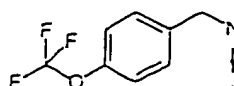
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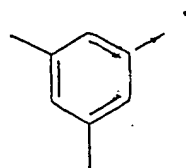
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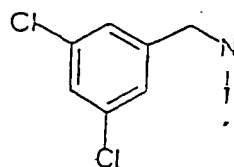
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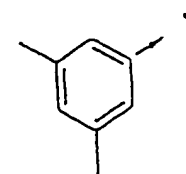
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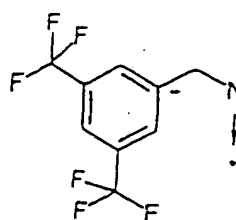
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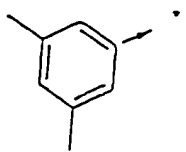
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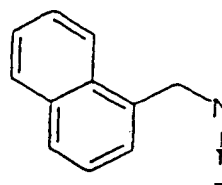
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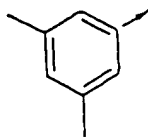
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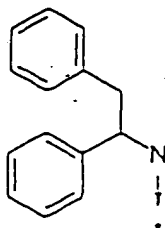
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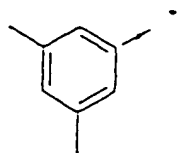
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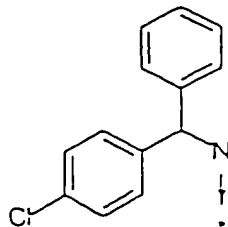
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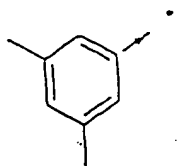
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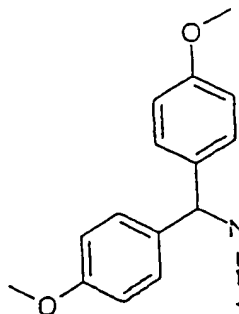
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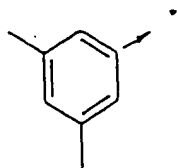
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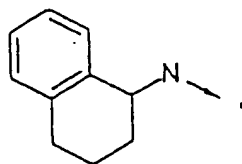
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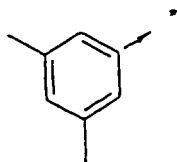
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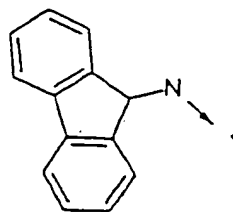
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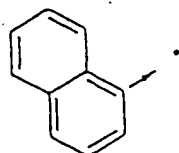
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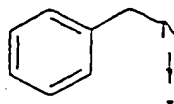
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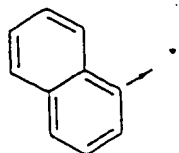
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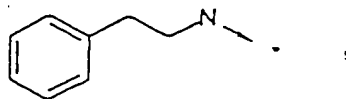
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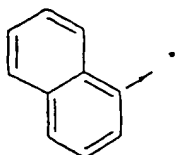
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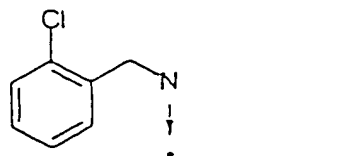
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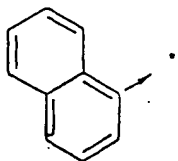
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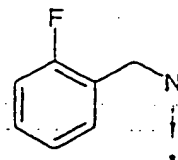
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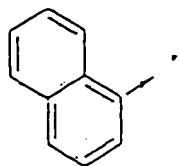
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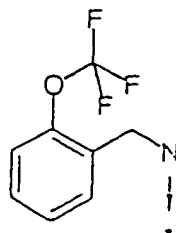
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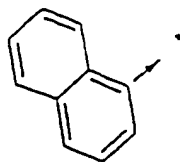
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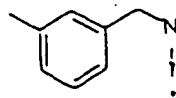
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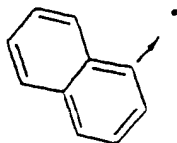
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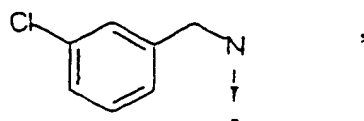
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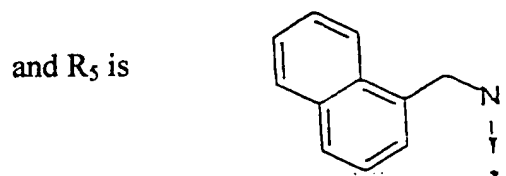
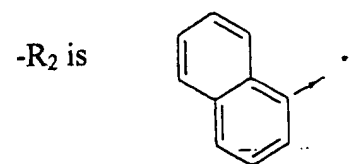
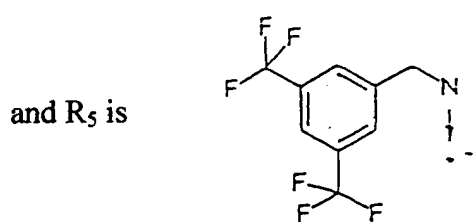
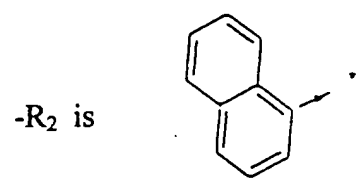
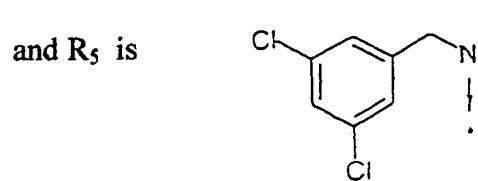
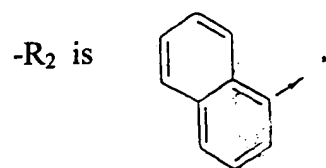
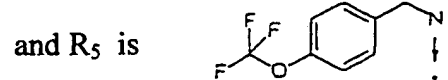
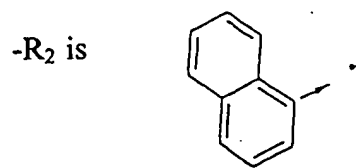
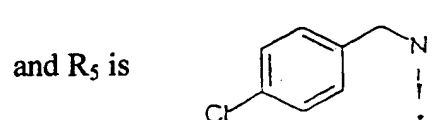
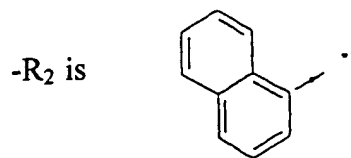
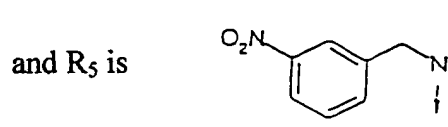
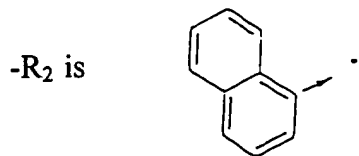
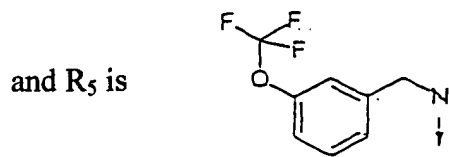
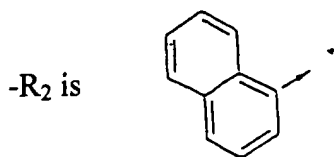


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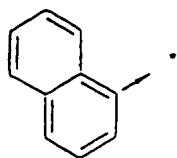


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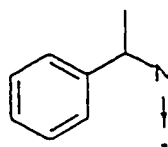




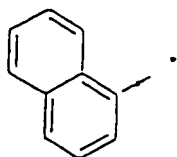
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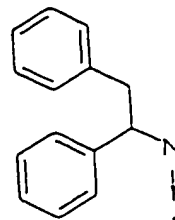
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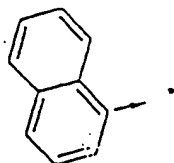
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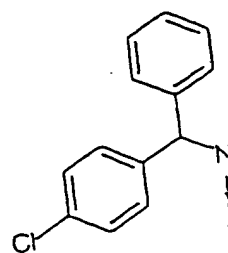
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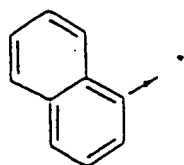
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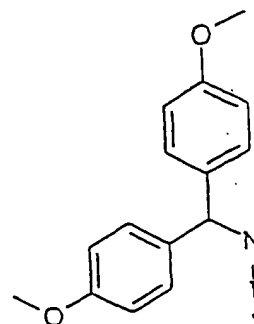
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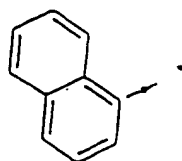
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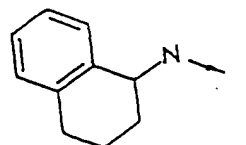


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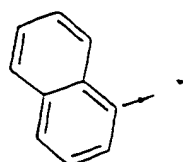


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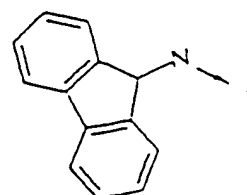
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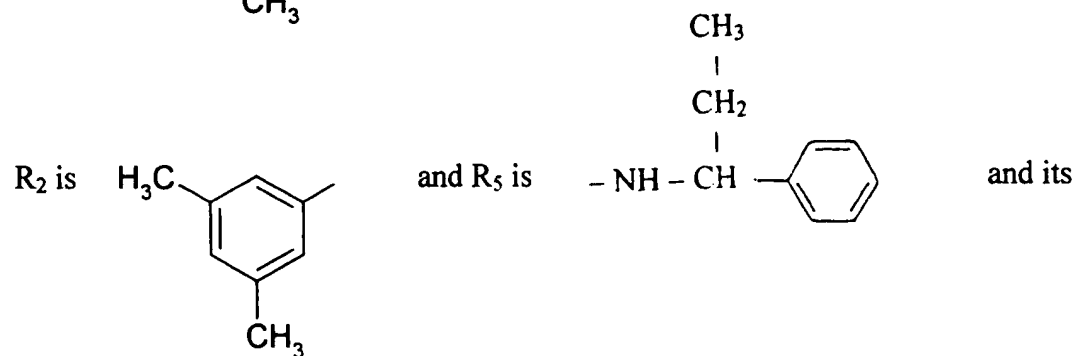
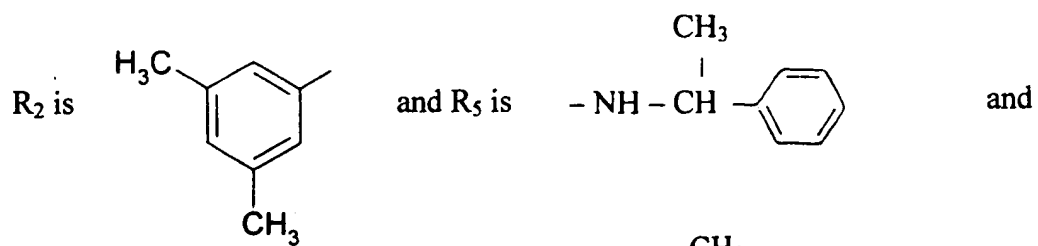


-R₂



and R₅ is





pharmaceutically acceptable salt.

